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JAN 17 2002

TECH CENTER 1600/2900

SEQUENCE LISTING

<110> Stanton, Jr., Vincent P.

<120> GENE SEQUENCE VARIANCE IN GENES RELATED
TO FOLATE METABOLISM HAVING UTILITY IN DETERMINING THE
TREATMENT OF DISEASE

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<212> DNA

<213> Homo sapiens

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<222> 166, 3432, 3682, 3937

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<222> 577, 638, 1708, 3730, 3925

<223> n = a or g

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<211> 2816

<212> DNA

<213> Homo sapiens

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 <222> 175, 1067
 <223> n = g or a

<221> misc_feature
 <222> 341
 <223> n = c or g

<221> misc_feature
 <222> 791, 1997, 2618, 2653
 <223> n = t or c

<221> misc_feature
 <222> 1337
 <223> n = c or a

<221> misc_feature
 <222> 2107
 <223> nucleotide in position 2107 is g, or absent

<221> misc_feature
 <222> 2583
 <223> n = t or g

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 <212> DNA
 <213> Homo sapiens

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 <222> 431, 441
 <223> n = a or g

<221> misc_feature
 <222> 498
 <223> n = c or t

<221> misc_feature
 <222> 579, 599
 <223> n = g or c

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 <222> 1066
 <223> n = t or c

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 <223> n = a or g

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<212> DNA
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<223> n = c or t

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<222> 452, 640
<223> n = a or g

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<222> 492, 625
<223> n = c or a

<221> misc_feature
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60

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<213> Homo sapiens

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<223> n = c or a

<221> misc_feature

<222> 716, 1293, 2401, 2429, 2618, 3083, 3125, 3635, 4256, 4898, 5062, 5167, 11069, 13298, 14479, 14730, 14796, 15344, 15450, 15503, 15590, 15840, 16149

<223> n = a or g

<221> misc_feature

<222> 732, 1379, 1590, 2488, 3212, 5006, 11238, 11422, 11686, 12598, 13171, 13645, 13782, 13806, 13813, 14586, 14788, 15042, 15546, 15770

<223> n = c or t

<221> misc_feature

<222> 1322, 1688

<223> n = c or g

<221> misc_feature

<222> 2594, 11293, 16199, 16203

<223> n = g or t

<221> misc_feature

<222> 3619

<223> n = a or t

<221> misc_feature

<222> 14547

<223> nucleotide in position 14547 is t, or absent

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 <213> Homo sapiens

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<221> misc_feature
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<221> misc_feature
 <222> 524
 <223> n = c or g

<221> misc_feature
 <222> 1399
 <223> n = t or a

<221> misc_feature
 <222> 1636, 1738, 2259
 <223> n = c or t

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60

120

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 <212> DNA
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 <222> 183, 1299
 <223> n = g or a

<221> misc_feature
 <222> 483
 <223> n = c or t

<221> misc_feature
 <222> 601

<223> n = g or c

<400> 10

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<210> 11

<211> 5847

<212> DNA

<213> Homo sapiens

<220>

<221> misc_feature

<222> 124, 3346, 5024, 5484, 5650

<223> n = c or t

<221> misc_feature

<222> 439, 1333, 1979, 2151, 2469, 2977, 4784, 5268, 5631, 5733

<223> n = g or a

<221> misc_feature

<222> 1045

<223> nucleotide at position 1045 is c, or absent

<221> misc_feature

<222> 1046

<223> nucleotide at position 1046 is t, or absent

<221> misc_feature

<222> 2636, 5287

<223> n = c or g

<221> misc_feature

<222> 3118

<223> n = g or t

<221> misc_feature

<222> 3257, 4053

<223> n = a or c

<221> misc_feature

<222> 5440

<223> n = t or a

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 <213> Homo sapiens

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 <223> n = c or t

<221> misc_feature
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<222> 1217, 1647, 2282
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